CORRECTED VERSION *

PCT

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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 4 :
A61K 9/22, 31/19, 31/20
A61K 31/23

A1

(11) International Publication Number: WO 87/ 03198

(43) International Publication Date: 4 June 1987 (04.06.87)

(21) International Application Number: PCT/SE86/00538

(22) International Filing Date: 24 November 1986 (24.11.86)

(31) Priority Application Number: 8505569-7

(32) Priority Date: 25 November 1985 (25.11.85)

(33) Priority Country: SE

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(81) Designated States: AT (European patent), AU, BB, BE (European patent), BG, BR, CF (OAPI patent), CG (OAPI patent), CH (European patent), CM (OAPI patent), DE (European patent), DK, FI, FR (European patent), GA (OAPI patent), GB (European patent), HU, IT (European patent), JP, KP, KR, LK, LU (European patent), MC, MG, ML (OAPI patent), MR (OAPI patent), MW, NL (European patent), NO, RO, SD, SE (European patent), SN (OAPI patent), SU, TD (OAPI patent), TG (OAPI patent), US.

Published
With international search report.

(54) Title: USE OF A HYDROPHOBIC SUBSTANCE FOR PREPARING AN ENTERIC PREPARATION FOR TREATMENT OF OBESITY AND AN ENTERIC PREPARATION

(57) Abstract

Use of a hydrophobic substance for preparing an enteric preparation for treatment of obesity. The enteric preparation is an anteric coated capsule, tablet or microcapsules, containing a hydrophobic substance in combination with an emulsifier. In a method for weight reduction, an effective weight reducing dosage of said preparation can be orally administered to a human.

* (Referred to in PCT Gazette No. 19/1987, Section II)

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Use of a hydrophobic substance for preparing an enteric preparation for treatment of obesity and an enteric preparation.

The present invention relates to the use of a hydrophobic substance for preparing an enteric preparation for treatment of obesity. The enteric preparation is a capsule, a tablet or microcapsules coated with a coating resistant to gastric juice which dissolves in ileum, the distsal portion of the small intestine. The invention also relates to a method for weight reduction, wherein said enteric preparation is orally administered in a weight reducing dosage to a human. In a further aspect the invention also relates to an enteric preparation containing specific hydrophobic substances in combination with an emulsifier.

One of the greatest health problems in modern times is over-weight, which is due to the fact that man has not been able to adapt himself to new circumstances requiring less energy but on the whole just as much of certain essential nutrients as before. A heavy overweight, obesity, is a serious medical problem which also leads to many complications.

Overweight is in addition not only a medical problem, but also a matter of well-being and personal satisfaction. It is well known that there is a great demand for effective methods of slimming.

Many drugs having an effect on the weight control in one way or another have been developed. Different investigations have shown that such drugs can bring about a weight loss of about 1-4 kg, but that this weight loss rarely is sustained beyond 8 weeks therapy. These drugs can be of use for therapy during a short period, but hardly offers any long term solution. In the long run fatness apparently can only be treated by changes of the fare and a weight reduction can therefore only be attained by a dietary change (S. Galloway et al, Postgraduate Medical Journal 60, 19-26, 1984).

For most overweight subjects it is hard to effect a weight reduction by slimming, that is a negative energy balance. In spite of different types of special diets it has turned out to be hard to reduce the body weight by means of slimming programs (Läkartidningen 80, no. 50, 4911-15, 1983).

It is generally known that the intake of food is controlled by changes in the composition of nutrients in the blood acting on centers in the hypothalamus, but also by gastrointestinal mechanisms which have still not been explained. It is however known, by means of animal tests, that an extension of the stomach brings about that the animal discontinues to eat, whereas the animal continues to eat for a long time if the food is continuously taken away from the stomach. It has also presently been demonstrated (N.W. Read et al, Gastroenterologi 86, 274-80, 1984) that the presence of unabsorbed food in ileum delays the gastric emptying. By this delay an increased absorption of the ingested food will be attained.

It has now surprisingly turned out that it is possible to effect a reduced food intake by bringing unabsorbed food and especially hydrofobic substances therein into contact with the distal part of the small intestine, that is ileum, where a physiologically mediated mechanism having this effect is started. Tests have shown that ileal infusion of a fat emulsion in connection with a meal brings about that a smaller amount of food is ingested than what should otherwise be the case. This physiologial effect is supposed to depend on that certain types of hydrofobic substances are interacting with specific ileal receptors and thereby induce satiety attended by a reduced food intake.

This finding has been transformed into enteric preparations which are simple and convenient to use for an overweight subject.

The invention relates to the use of a hydrophobic substance for preparing an enteric preparation. In said preparation the hydrophobic substance is used in combination with an emulsifier.

The hydrophobic substance can be a fatty acid having 6-28 carbon atoms, an ester or a salt thereof, a fatty alcohol having 6-28 carbon atoms or an ester thereof.

The fatty acid can be saturated or unsaturated, and have a branched or a straight chain. As examples of fatty acids can be mentioned lauric acid, palmitic acid, stearic acid, oleic acid, ricinoleic acid, linoleic acid and linolenic acid. Preferred acids have 14-22 carbon atoms. Esters of fatty acids can be esters of one or more of said acids and an alcohol containing one ore more hydroxyl groups. In a diester one of the ester groups can be derived from a fatty acid having 2-5 carbon atoms. The alcohol can have 2-28 carbon atoms. Examples of esters are isopropyl isostearate, sorbitan trioleate (Span® 85), 1,2-propylene glycol myristate, dipalmitin, acetylated monoglyceride from soybean oil, glyceryl palmitate lactate, hydrogenated tallow glycerides citrate, diacetyl tartaric acid esters of monooleate. A salt refers to a physiologically acceptable salt of an inorganic ion, e.g. sodium, potassium, magnesium or calcium, or an ammonium, mono-, di- or triethanolammonium ion.

The fatty alcohol having 6-28 carbon atoms can be saturated or unsaturated and have a branched or a straight chain. An ester of a fatty alcohol can be an ester of an acid having 1-28 carbon atoms. As an example can be mentioned arachidyl propionate.

As examples of fats, that is mixtures of esters of fatty acids of 6-28 carbon atoms and glycerol, can be mentioned vegetable and animal fats and oils such as borage oil, corn oil, sesame

oil, olive oil, soybean oil, cottonseed oil, castor oil, peanut oil, cocoa butter, cod-liver oil. Said mixtures can optionally also contain other substances, such as fatty acids and fatty alcohols.

The hydrophobic substance in the enteric preparation can be in liquid form, such as an oil or a solution or an emulsion of a solid or liquid substance. The hydrophobic substances can also be solid at ambient temperature, provided they are fluid in the intestine.

As emulsifier or dispersing agent can be used tensides and amphoteric compounds which are common within pharmacy, for instance Tween^{R} 80 (polyoxyethylene glycol sorbitan mono-oleat), Arlaton^{R} T (polyoxyethylene fatty acid ester), sodium taurocholate, albumin and lecithin, which are described in Martindale, The Extrapharmacopoeia, 28th ed., 1982. The emulsifier is combined with the hydrophobic substance in an amount of at least 1% by weight based on the hydrophobic substance. The amount to be added should be sufficient to provide a spontaneous emulgation of the hydrophobic substance in the intestine. Free fatty acids generally do not require as much emulsifier as for instance triglycerides of fatty acids. More than 30% should however not be added, a range of 3-15% being preferable.

The enteric prepartion is a capsule, a tablet or micro-capsules containing the hydrophobic substance optionally in combination with the emulsifier, which has been coated with a gastric juice resistant coating. The capsules can be hard or soft gelatin capsules containing the active substance in solid and preferably liquid form respectively. Said capsules can also be designed for a slow release of the active substance. Tablets are preferably prepared by moulding the hydrophobic substance, optionally with additives, in accordance with conventional

technique. In order to make the preparations unsolvable in acid medium, that is resistant to gastric juice, they are finally coated with a coating of so called gastric juice resistant polymers, which dissolve in the distal portion of the small intestine where the actual receptors are located. As examples of suitable polymers which can be used as coatings can be mentioned cellulose derivatives esterified with phthalic acid, such as cellulose acetate phthalate (CAP), hydroxypropylmethyl-cellulose phthalate (HP 50, HP 55), and polymers based on methacrylic acid (Eudragit^{R} L, Röhm Pharma GmbH) and copolymers based on methacrylic acid methylester (Eudragit^{R} S, Röhm Pharma GmbH). These polymers are applied, if required, in admixture with plasticizers and other conventional additives, for instance pigments. It should of course be seen to it that the polymer coating is designed in such a way that it dissolves and releases its content at the intended place in the intestine. Microcapsules can be manufactured by emulsifying the hydrophobic substance in a water solution of a polymer, e.g. cellulose acetate phthalate, and precipitation of the polymer, that is by coacervation, or by spray coating, also in accordance with conventional technique.

The invention also relates to a method for weight reduction, wherein an effective weight reducing dosage of an enteric preparation containing a hydrophobic substance is orally administered to a human. The enteric preparation should be taken in good time before each meal and in order to achieve optimal effect the hydrophobic substance in the preparation should have been brought into contact with the actual receptors in ileum before the meal is started. This means that one or more capsules or tablets or microcapsules containing 1-5 g hydrophobic substance and emulsifier should be taken starting 2-5 hours, preferably 2-3 hours before each meal from 1 to 6 times/d. It is of advantage if the preparation is formulated in such a way that the hydrofobic substance is released for a longer period. One dose unit normally comprises 0.01-1 g of the mixture of the hydrophobic substance and the emulsifier.

The invention also relates to an enteric preparation, which is a capsule, a tablet or microcapsules, containing a pharmaceutically active substance and being coated with a gastric juice resistant coating, in which the active substance is a saturated or unsaturated fatty acid having 6-28 carbon atoms, a physiologically acceptable salt thereof, a triglyceride of one or more saturated or unsaturated fatty acids having 6-22 carbon atoms, or a mixture thereof, which is combined with 1-30% by weight based on the hydrophobic substance of an emulsifier.

In a preferred preparation the active substance is a saturated or unsaturated fatty acid having 14-22 carbon atoms, a physiologically acceptable salt thereof or a mixture therof, which is combined with 3-15% by weight based on the hydrophobic substance of an emulsifier.

The weight reducing effect that is attained by the administration of a hydrophobic substance to ileum will be apparent from the following test.

Infusion of fat emulsion to humans

A fat emulsion comprising 50% corn oil and 3% albumin, and a control solution which only contained 3% albumin were administrated separately at an interval of about a week by ileal infusion to a group of 6 persons in connection with the eating of a appetizing meal.

As a comparison a 20% fat emulsion, essentially containing fractionated soybean oil, Intralipid® (Kabi Vitrum), and a control solution of 0.9% saline were administrated by intravenous infusion to another group of 6 persons under otherwise like conditions.

All the test persons ate until they experienced a feeling of fullness which was scored in the same way on a scale from 1 to 10. The time taken to eat the meal and the quantity of ingested food and drink, as well as the caloric intake calculated therefrom is evident from the table below.

Table 1

The effect of ileal or intravenous infusion of fat emulsion on food intake

	Ileal infusion		Intravenous infusion	
	Control	Fat emulsion	Control	Fat emulsion
Duration of the meal (min)	31.7±3.0	24.8±1.0	27.1±2.8	25.5±2.6
Amount of food intake (g)	884±89	670±23	902±39	934±109
Amount of drink intake (ml)	490±53	447.5±46	550±62	558±61
Total energy intake (kcal)	1883±259	1313±90	1965±137	1938±236

It follows that an ileal infusion of corn oil reduces the amount of food eaten and also the total energy intake. This reduction was greater than the amount of energy which was supplied by the infusion of the fat emulsion. This difference in food intake is not obtained in intravenous infusion of a fat emulsion.

Rat infusion test

The effect of different hydrophobic substances on the gastric emptying was evaluated by means of the following paired study on rats, in which the transit time for an unabsorbable carbohydrate from stomach to caecum was measured by means of hydrogen analysis (see N.J.Brown et al, "Adaption of hydrogen analysis to measure stomach to caesum transit time in the rat", Department of Physiology, The University, Western Bank, Sheffield).

Male albino rats, weighing 250-300 g each, which had been fasted for 18 hours were, after infusion of a test solution or control for 20 minutes through an abdominal sond to the ileum

gavaged with 5 ml homogenized baked beans and 0.5 g lactose. The transit of the head of said meal from the stomach to the caecum was assessed by the rise in hydrogen in the environment of the rat, deriving from the fermentation of the unabsorbable carbohydrates in the meal by colon bacteria. The transit time for the test solution and the control were recorded.

The hydrophobic substances to be tested were dispersed in water or emulsified in saline by means of an emulsifier to a concentration of 10-20% by weight. Tween® 80, egg lecithin and/or sodium taurocholate were added to the substances as emulsifiers in an amount of 2-12% by weight based on the hydrophobic substance. The control solution was 0.9% saline to which emulsifier had been added in an amount corresponding to the test solution content. The results of the test are shown in Table 2.

Table 2

Influence of hydrophobic substances on stomach to caecum transit time

Hydrophobic substance	Concentration %	Transit time * min
Distilled rapeseed fatty acids **, ***	20	210
Borage oil **	20	190
Soybean oil + 10% distilled rapeseed fatty acids **,***	20	138
Corn oil + 10% distilled rapeseed fatty acids **,***	20	147
Control	-	107

* average of 2-4 values
** containing 6.3% egg lecithin and 2.5% sodium taurocholate
*** containing 3.5% Tween® 80

It is believed that there is a relation between reduced food intake and delay in gastric emptying.

Example 1

A tablet of the invention can be prepared by moulding a fat which is solid at ambient temperature, such as cocoa butter, in accordance with common technique. Tablets prepared in this way are then coated by means of a coating equipment. For the coating of 5 kg tablets of nonstop type the following composition can for example be used.

Coating composition	Amount, g
polyethylene glycol 6000	10.0 g
propylene glycol	39.0 g
sorbitan oleat	13.0 g
cellulosaacetate phthalate	130 g
ethanol 70%	717 g
acetone	1091 g

After the coating micronised talc is applied 2-5 times in a total amount of 125 g. Tablets coated in this way will obtain a gastric juice resistant coating which is dissolved in ileum.

Example 2

An enteric capsule for oral administration of a hydrophobic substance can be manufactured by loading a soft gelatine capsule (size No. 10 oval, from R.P. Scherer Corporation) with the following mixture.

Capsule content	% by weight	Amount, mg
corn oil	87	435
distilled rapeseed fatty acids	10	50
Tween® 80	3	15
	in total	500

After filling and sealing by means of conventional technique, the capsule was spraycoated with the following solution.

Coating composition	Approximate amount, mg
hydroxypropylmethylcellulose phthalat (HP 55)	32
dibutylic phthalate	6.4
ethanol containing 0.75% methylethylketone	400
water	70

The spraycoating is continued until resistance to gastric juice is obtained (according to USP, Pharm. Eur.).
The weight of the coating is then 38.5 mg.

C L A I M S

1. Use of a hydrophobic substance for preparing an enteric preparation for treatment of obesity.

2. Use according to claim 1, characterized in that the hydrophobic substance is used in combination with an emulsifier.

3. Use according to claim 1 or 2, characterized in that the enteric preparation is a capsule, a tablet or microcapsules, coated with a gastric juice resistant coating.

4. Use according to any of claims 1-3, characterized in that the hydrophobic substance is a saturated or unsaturated fatty acid having 6-28 carbon atoms, a physiologically acceptable salt thereof or an ester of one or more of said acids and an alcohol containing one or more hydroxyl groups.

5. Use according to any of claims 1-4, characterized in that hydrophobic substance is a glycerol ester of one or more saturated or unsaturated fatty acids having 14-22 carbon atoms or a mixture thereof.

6. Use according to any of claims 1-4, characterized in that that the hydrophobic substance is a saturated or unsaturated fatty acid having 14-22 carbon atoms or a physiologically acceptable salt thereof, or a mixture thereof.

7. Method for weight reduction, characterized in orally administering an effective weight reducing dosage of an enteric preparation containing a hydrophobic substance to a human.

8. Method according to claim 7, wherein the enteric preparation is a capsule, a tablet or microcapsules coated with a gastric juice resistant coating, containing 0.01-1 g of a mixture of the hydrophobic substance and an emulsifier, characterized in that 1-5 g of said mixture is administrated 2-3 h before every meal.

9. Enteric preparation, which is a capsule, a tablet or microcapsules containing a pharmaceutically active substance and being coated with a gastric juice resistant coating, characterized in that the active substance is a saturated or unsaturated fatty acid having 6-28 carbon atoms, a physiologically acceptable salt thereof, a triglyceide of one or more saturated or unsaturated fatty acids having 6-22 carbon atoms, or a mixture thereof, which is combined with 1-30% by weight, based on the hydrophobic substance, of an emulsifier.

10. Enteric preparation according to claim 9, characterized in that the active substance is a saturated or unsaturated fatty acid having 14-22 carbon atoms, a physiologically acceptable salt thereof or a mixture thereof, which is combined with 3-15% by weight, based on the hydrophobic substance, of an emulsifier.

INTERNATIONAL SEARCH REPORT

International Application No PCT/SE86/00538

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all) [6]

According to International Patent Classification (IPC) or to both National Classification and IPC [4]

A 61 K 9/22, 31/19, /20, /23

II. FIELDS SEARCHED

Minimum Documentation Searched [7]

Classification System	Classification Symbols
IPC	A 61 K 9/00, /22, /24, /36, /42, /52, /62, 31//19, /20, /23, /25
US Cl	424: 16, 19, 23, 24, 35, 38

Documentation Searched other than Minimum Documentation to the Extent that such Documents are included in the Fields Searched [8]

SE, NO, DK, FI classes as above

III. DOCUMENTS CONSIDERED TO BE RELEVANT [9]

Category [10]	Citation of Document, [11] with indication, where appropriate, of the relevant passages [12]	Relevant to Claim No. [13]
Y	DE, A, 2 344 958 (ZAHN H) 20 March 1975 see claims	1,4,6
Y	EP A, 133 110 (SANOFI) 13 February 1985 see page 2, lines 1-3 and 13-21 and page 7, lines 1-8 and claims & AU, 30755/84 FR, 2549370 JP, 60041610 FR, 2549374	1,4
A	GB, A, 2 140 688 (OTSUKA PHARMA CEUTICAL & CO) 5 December 1984 see claims	
A	GB, A, 1 142 804 (LABORATORIES PHARMASCIENCE) 12 February 1969 see claims	

* Special categories of cited documents: [10]

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

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"P" document published prior to the international filing date but later than the priority date claimed

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"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

IV. CERTIFICATION

Date of the Actual Completion of the International Search	Date of Mailing of this International Search Report
1987-01-12	1987 -01- 27
International Searching Authority	**Signature of Authorized Officer**
Swedish Patent Office	Agneta Tannerfeldt

MJ

Form PCT/ISA/210 (second sheet) (January 1985)

FURTHER INFORMATION CONTINUED FROM THE SECOND SHEET

V. [X] OBSERVATIONS WHERE CERTAIN CLAIMS WERE FOUND UNSEARCHABLE [1]

This International search report has not been established in respect of certain claims under Article 17(2) (a) for the following reasons:

1. [X] Claim numbers 7-8, because they relate to subject matter not required to be searched by this Authority, namely:

Method for treatment of the human or animal body.

2. [ ] Claim numbers....., because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

3. [ ] Claim numbers....., because they are dependent claims and are not drafted in accordance with the second and third sentences of PCT Rule 6.4(a).

VI. [ ] OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING [2]

This International Searching Authority found multiple inventions in this international application as follows:

1. [ ] As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims of the international application.

2. [ ] As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims of the international application for which fees were paid, specifically claims:

3. [ ] No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claim numbers:

4. [ ] As all searchable claims could be searched without effort justifying an additional fee, the International Searching Authority did not invite payment of any additional fee.

Remark on Protest

[ ] The additional search fees were accompanied by applicant's protest.

[ ] No protest accompanied the payment of additional search fees.

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Use of a hydrophobic substance for preparing an enteric preparation for treatment of obesity and an enteric preparation.

The present invention relates to an enteric preparation, which induces satiety and thereby a reduced food intake. The enteric preparation is a capsule or a tablet containing a pharmaceutically active substance, and which is coated with a coating resistant to gastric juice which dissolves in ileum, the distal portion of the small intestine. The invention also relates to the use of said preparation for effecting a weight reduction in an overweight subject.

One of the greatest health problems in modern times is overweight, which is due to the fact that man has not been able to adapt himself to new circumstances requiring less energy but on the whole just as much of certain essential nutrients as before. A heavy overweight, obesity, is a serious medical problem which also leads to many complications.

Many drugs having an effect on the weight control in one way or another have been developed. Different investigations have shown that such drugs can bring about a weight loss of about 1-4 kg, but that this weight loss rarely is sustained beyond 8 weeks therapy. These drugs can be of use for therapy during a short period, but hardly offers any long term solution. In the long run fatness apparently can only be treated by changes of the fare and a weight reduction can therefore only be attained by a dietary change (S. Galloway et al, Postgraduate Medical Journal 60, 19-26, 1984).

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It is generally known that the intake of food is controlled by changes in the composition of nutrients in the blood acting on centers in the hypothalamus, but also by gastrointestinal mechanisms which have still not been explained. It is however known, by means of animal tests, that an extention of the stomach brings about that the animal discontinues to eat, whereas the animal continues to eat for a long time if the food is continuously taken away from the stomach. It has also presently been demonstrated (N.W. Read et al, Gastroenterologi 86, 274-80, 1984) that the presence of unabsorbed food in ileum delays the gastric emptying. By this delay an increased absorption of the ingested food will be attained.

It has now surprisingly turned out that it is possible to effect a reduced food intake by bringing unabsorbed food and especially hydrofobic substances therein into contact with the distal part of the small intestine, that is ileum, where they start a physiologically mediated mechanism having this effect. Tests have shown that ileal infusion of a fat emulsion in connection with a meal brings about that a smaller amount of food is ingested than what should otherwise be the case. This physiologial effect is supposed to depend on that certain types of hydrofobic substances are interacting with specific ileal receptors and thereby induce satiety attended by a reduced food intake.

This finding has than been transformed into pharmaceutical compositions which are simple and convenient to use for an overweight subject.

The invention thus relates to an enteric preparation in the form of a capsule or a tablet coated with a coating resistant to gastric juice which is dissolved in ileum, which is characterized in containing, as an active substance, a hydrofobic substance interfering with specific receptors in ileum and then inducing satiety and reduced food intake.

The hydrofobic substance is for instance a lipid, such as a fat or an oil, a phosphatide or a wax. Suitable lipids are a saturated or unsaturated fatty acid having 6-28 carbon atoms, ester or an ether of such a fatty acid and a mono or multivalent alcohol, a salt of such a fatty acid and an amino compound, or a mixture of such compounds. Among saturated and unsaturated fatty acids of 6-28 carbon atoms, polyunsaturated acids included, can be mentioned butyric acid, lauric acid, palmitic acid, stearic acid, oleic acid, ricinoleic acid, linoleic acid and linolenic acid. Esters and ethers of said fatty acids are for instance those with glycerol, that is mono-, di- and triacylglycerol and mono-, di- and trialkylglycerol. A polyhydric alcohol that can form an ester or an ether with said fatty acids is for instance a carbohydrate.

As examples of fats, that is mixtures of esters of fatty acids of 6-28 carbon atoms and glycerol, can be mentioned vegetable and animal fats and oils such as corn oil, sesame oil, olive oil, soybean oil, cottonseed oil, castor oil, peanut oil, cocoa butter, cod-liver oil. Said mixtures can optionally also contain other substances, such as fatty acids and fatty alcohols. The hydrofobic substance in the enteric preparation can be in liquid form, such as an oil or a solution or an emulsion of a solid or liquid substance. The hydrofobic substance can also be solid at ambient temperature. As emulsifier or dispersing agent can be used tensides and amphoteric compounds which are common in the pharmacy, for instance Tween® 80 (polyoxyethylene glycol sorbitan mono-oleat), Arlaton® T (polyoxyethylene fatty acid ester), albumin and lecithin, which are described in Martindale, The Extrapharmacopoeia, 28th ed., 1982.

The enteric prepartion is a capsule or a tablet containing the active substance, which has been coated with a gastric juice resistant coating. The capsules can be hard or soft gelatin

capsules containing the active substance in solid and liquid form respectively. Said capsules can also be designed for a slow release of the active substance. Tablets are preferably prepared by moulding the hydrofobic substance, optionally with additives, in accordance with conventional technique. In order to make the capsules and tablets unsolvable in acid medium, that is resistant to gastric juice, they are finally coated with a coating of so called gastric juice resistant polymers, which dissolve in the distal portion of the small intestine where the actual receptors are located. As examples of suitable polymers which can be used as coatings can be mentioned cellulose derivatives esterified with phthalic acid, such as cellulose acetate phthalate (CAP), hydroxypropylmetnyl-cellulose phthalate (HP 50, HP 55), and polymers based on methacrylic acid (Eudragit® L, Röhm Pharma GmbH) and copolymers based on methacrylic acid methylester (Eudragit® S, Röhm Pharma GmbH). These polymers are applied, if required, in admixture with softeners and other conventional additives, for instance pigments. It should of course be seen to it that the polymer coating is designed in such a way that it dissolves and releases its content at the intended place in the intestine.

The invention also relates to the use of an enteric preparation being a capsule or a tablet having a gastric juice resistant coating which is dissolved in ileum, which contains a hydrofobic substance interfering with specific receptors i ileum and then inducing satiety and reduced food intake as an active substance, in order to effect weight reduction in an overweight subject. The enteric preparation should be taken in good time before each meal and in order to achieve optimal effect the hydrofobic substance in the preparation should have been brought into contact with the actual receptors in ileum before the meal is started. This means that one or more capsules or tablets should be taken starting 2-5 hours, preferably

2-3 hours before each meal. It is of advantage if the preparation is formulated in such a way that the hydrofobic substance is released for a longer period. One dose unit normally comprises 0.2-1.2 ml of a liquid hydrofobic substance or 0.1-1.5 g if the hydrofobic substance is solid.

The weight reducing effect that is attained by the administration of a hydrofobic substance to ileum will be apparent from the following test.

INFUSION OF FAT EMULSION

A fat emulsion comprising 50% corn oil and 3% albumin, and a control solution which only contained 3% albumin were administrated separately at an interval of about a week by ileal infusion to a group of 6 persons in connection with the eating of a appetizing meal.

As a comparison a 20% fat emulsion, essentially containing fractionated soybean oil, Intralipid® (Kabi Vitrum), and a control solution of 0.9% saline were administrated by intravenous infusion to another group of 6 persons under otherwise like conditions.

All the test persons ate until they experienced a feeling of fullness which was scored in the same way on a scale from 1 to 10. The time taken to eat the meal and the quantity of ingested food and drink, as well as the caloric intake calculated therefrom is evident from the table below.

TABLE

The effect of ileal or intravenous infusion of fat emulsion on food intake

	Ileal infusion		Intravenous infusion	
	Control	Fat emulsion	Control	Fat emulsion
Duration of the meal (min)	31.7±3.0	24.8±1.0	27.1±2.8	25.5±2.6
Amount of food intake (g)	884±89	670±23	902±39	934±109
Amount of drink intake (ml)	490±53	447.5±46	550±62	558±61
Total energy intake (kcal)	1883±259	1313±90	1965±137	1938±236

It follows that an ileal infusion of corn oil reduces the amount of food eaten and also the total energy intake. This reduction was greater than the amount of energy which was supplied by the infusion of the fat emulsion. This difference in food intake is not obtained in intravenous infusion of a fat emulsion.

A tablet of the invention can be prepared by moulding a fat which is solid at ambient temperature, such as cocoa butter, in accordance with common technique. Tablets prepared in this way are then coated by means of a coating equipment. For the coating of 5 kg tablets of nonstoptype the following composition can for example be used.

COATING COMPOSITION

polyethylene glycol 6000	10.0 g
propylene glycol	39.0 g
sorbitan oleat	13.0 g
celluloseacetate phthalate	130 g
ethanol 70%	717 g
acetone	1091 g

After the coating micronised talc is applied 2-5 times in a total amount of 125 g. Tablets coated in this way will obtain a gastric juice resistant coating which is dissolved in ileum.

PATENT CLAIMS

1. Enteric preparation, which is a capsule or a tablet containing a pharmaceutically active substance and being coated with a gastric juice resistant coating which dissolves in ileum, c h a r a c t e r i z e d i n that the active substance is a hydrofobic substance interfering with specific receptors in ileum and then inducing satiety and a reduced food intake.

2. Enteric preparation according to claim 1, c h a r a c t e r i z e d i n that the hydrofobic substance is a lipid.

3. Enteric preparation according to claim 1, c h a r a c t e r i z e d i n that the hydrofobic substance is a saturated or unsaturated fatty acid having 6-28 carbon atoms, an ester or an ether of such a fatty acid and a mono or polyhydric alcohol, a salt of such a fatty acid and an amino compound, or a mixture thereof.

4. Enteric preparation according to any of the claims 1-3, c h a r a c t e r i z e d i n that the hydrofobic substance is a mixture of esters of a fatty acid having 6-28 carbon atoms and glycerol.

5. Enteric preparation according to any of the claims 1-4, c h a r a c t e r i z e d i n that the gastric juice resistant coating which is dissolved in ileum is a cellulose derivative esterified with phthalic acid or a polymer based on methacrylic acid or methacrylic acid methylester, or a mixture thereof, if required together with softeners and other conventional additives.

6. Enteric preparation according to any of the claims 1-5, characterized in that it is a soft gelatin capsule, which contains a liquid fat or a solution or an emulsion of a solid or liquid fat, and which is coated with a coating of hydroxipropylmethylcellulose phthalate.

7. Use of an enteric preparation for effecting a weight reduction in an overweight subject, which preparation is a capsule or a tablet having a gastric juice resistant coating, which dissolves in ileum, which contains, as an active substance, a hydrofobic substance interfering with specific receptors in ileum and then inducing satiety and a reduced food intake.

INTERNATIONAL SEARCH REPORT

International Application No PCT/SE86/00538

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all) [6]

According to International Patent Classification (IPC) or to both National Classification and IPC [4]

A 61 K 9/22, 31/19, /20, /23

II. FIELDS SEARCHED

Minimum Documentation Searched [7]

Classification System	Classification Symbols
IPC	A 61 K 9/00, /22, /24, /36, /42, /52, /62, 31//19, /20, /23, /25
US Cl	424: 16, 19, 23, 24, 35, 38

Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched [8]

SE, NO, DK, FI classes as above

III. DOCUMENTS CONSIDERED TO BE RELEVANT [9]

Category [10]	Citation of Document, [11] with indication, where appropriate, of the relevant passages [12]	Relevant to Claim No. [13]
Y	DE, A, 2 344 958 (ZAHN H) 20 March 1975 see claims	1,4,6
Y	EP A, 133 110 (SANOFI) 13 February 1985 see page 2, lines 1-3 and 13-21 and page 7, lines 1-8 and claims & AU, 30755/84 FR, 2549370 JP, 60041610 FR, 2549374	1,4
A	GB, A, 2 140 688 (OTSUKA PHARMA CEUTICAL & CO) 5 December 1984 see claims	
A	GB, A, 1 142 804 (LABORATORIES PHARMASCIENCE) 12 February 1969 see claims	

* Special categories of cited documents: [10]

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

IV. CERTIFICATION

Date of the Actual Completion of the International Search	Date of Mailing of this International Search Report
1987-01-12	1987 -01- 27
International Searching Authority	**Signature of Authorized Officer**
Swedish Patent Office	Agneta Tannerfeldt

MJ

Form PCT/ISA/210 (second sheet) (January 1985)

FURTHER INFORMATION CONTINUED FROM THE SECOND SHEET

V. [X] OBSERVATIONS WHERE CERTAIN CLAIMS WERE FOUND UNSEARCHABLE [1]

This international search report has not been established in respect of certain claims under Article 17(2) (a) for the following reasons:

1. [X] Claim numbers 7-8, because they relate to subject matter not required to be searched by this Authority, namely:

Method for treatment of the human or animal body.

2. [ ] Claim numbers....., because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

3. [ ] Claim numbers....., because they are dependent claims and are not drafted in accordance with the second and third sentences of PCT Rule 6.4(a).

VI. [ ] OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING [2]

This International Searching Authority found multiple inventions in this international application as follows:

1. [ ] As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims of the international application.
2. [ ] As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims of the international application for which fees were paid, specifically claims:
3. [ ] No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claim numbers:
4. [ ] As all searchable claims could be searched without effort justifying an additional fee, the International Searching Authority did not invite payment of any additional fee.

Remark on Protest

- [ ] The additional search fees were accompanied by applicant's protest.
- [ ] No protest accompanied the payment of additional search fees.